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FILE LAST UPDATED: 23 May 2008 (20080523/ED)

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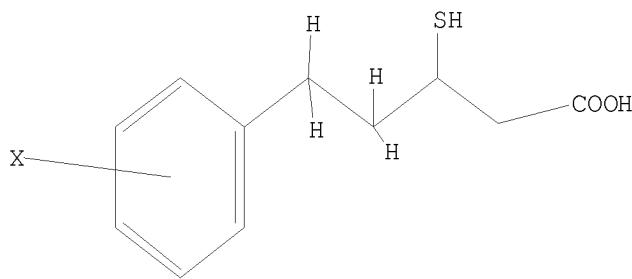
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L1 STRUCTURE UPLOADED

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L1 HAS NO ANSWERS
L1 STR

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Structure attributes must be viewed using STN Express query preparation.

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REG1stRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
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FULL SEARCH INITIATED 17:25:52 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 2000 TO ITERATE

100.0% PROCESSED 2000 ITERATIONS 2 ANSWERS
SEARCH TIME: 00.00.01

L2 2 SEA SSS FUL L1

L3 1 L2

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L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:1006952 CAPLUS
DOCUMENT NUMBER: 140:59517
TITLE: Preparation of 2,5-disubstituted 3-mercaptopentanoic acids as carboxypeptidase U inhibitors
INVENTOR(S): Polla, Magnus
PATENT ASSIGNEE(S): AstraZeneca A.B., Swed.
SOURCE: PCT Int. Appl., 38 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2003106420	A1	20031224	WO 2003-SE970	20030610

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
 CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
 GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
 LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
 PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT,
 TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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 CA 2488606 A1 20031224 CA 2003-2488606 20030610
 AU 2003241260 A1 20031231 AU 2003-241260 20030610
 AU 2003241260 B2 20070426
 BR 2003011384 A 20050315 BR 2003-11384 20030610
 EP 1532111 A1 20050525 EP 2003-730987 20030610
 EP 1532111 B1 20070808
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 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
 CN 1662504 A 20050831 CN 2003-813840 20030610
 JP 2005533064 T 20051104 JP 2004-513253 20030610
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 MX 2004PA12604 A 20050323 MX 2004-PA12604 20041214
 HK 1077296 A1 20071109 HK 2005-109105 20051014
 PRIORITY APPLN. INFO.: SE 2002-1837 A 20020614
 WO 2003-SE970 W 20030610

OTHER SOURCE(S): MARPAT 140:59517

AB The title compds. R1(CH2)2CH(SH)CH(CO2H)CH2R2 [I; R1 = (un)substituted Ph, naphthyl, pyridinyl, etc.; R2 = aminopyridinyl, aminothiazolyl, 3-azabicyclo[3.2.1]octyl] which inhibit carboxypeptidase U and thus can be used in the prevention and treatment of diseases where inhibition of carboxypeptidase U is beneficial, were prepared E.g., a 4-step synthesis of 2-[(6-aminopyridin-3-yl)methyl]-5-(1,1'-biphenyl-3-yl)-3-mercaptopentanoic acid (starting from 3-iodo-1,1'-biphenyl), was given. Biol. data was given for 13 exemplified compds. I. In further aspects, the invention relates to pharmaceutical compns. containing at least one compound I.

IT 637300-48-2P 637300-49-3P

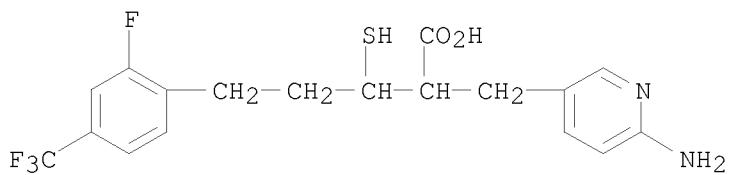
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 2,5-disubstituted 3-mercaptopentanoic acids as carboxypeptidase U inhibitors)

RN 637300-48-2 CAPLUS

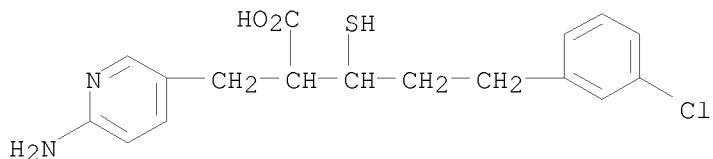
CN 3-Pyridinepropanoic acid, 6-amino- α -[3-[2-fluoro-4-(trifluoromethyl)phenyl]-1-mercaptopropyl]- (CA INDEX NAME)

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RN 637300-49-3 CAPLUS

CN 3-Pyridinepropanoic acid, 6-amino- α -[3-(3-chlorophenyl)-1-mercaptopropyl]- (CA INDEX NAME)



REFERENCE COUNT:

1

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